WEST Search History

DATE: Monday, August 18, 2003

<u>Set</u> <u>Name</u> side by side	Query	<u>Hit</u> Count	Set Name result set
DB=US	PT; PLUR=YES; OP=AND		
L1	igy.clm. and campylobact\$.clm.	0	L1
L2	ig-y.clm. and campylobact\$.clm.	0	L2
L3	campylobact\$.clm. or jejuni.clm.	191	L3
L4	L3 same (avian or turkey or fowl or poultry or chick or chicken or bird)	7	L4
L5	anticampylobacter.clm. or anti-campylobacter.clm.	0	L5
L6	anti-Campylobacter.clm.	3	L6
L7	(campylobacter or jejuni) same (igy or igg or igm or iga or antisera or antiserum or anti-sera or anti-serum or immune or immunoglobulin or monoclonal or polyclonal or poly-clonal or mono-clonal)	123	L7
L8	L7 same (avian or turkey or fowl or poultry or chick or chicken or bird)	16	L8
L9	campylobact\$.clm. or jejuni.clm.	191	L9
L10	L9 and (flge or flg-e or hook).clm.	0	L10
L11	(campylobact\$ or jejuni).ti,ab. and hook.clm.	0	L11
L12	(campylobact\$ or jejuni).ti,ab. and flgE.clm.	0	L12
L13	(campylobact\$ or jejuni) and flgE	2	L13

END OF SEARCH HISTORY

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(S1 OR S2 OR S3) AND (
                                         AN (5N) (ANTIBOD? OR ANTISER?
S4
             IMMUNE? OR SEROTYP? OR IMMUNE? OR IMMUNOGLOB? OR IGA OR IGM OR
              IGG?))
S5
        12196
                (AVIAN OR CHICK? OR FOWL? OR TURKEY? OR BIRD? OR IGY?) (5N)
              (ANTIBOD? OR ANTISER? OR IMMUNE? OR SEROTYP? OR IMMUNE? OR I-
                                                                            2/03
MACOS
             MMUNOGLOB? OR IGA OR IGM OR IGG?)
S6
           45
                (S1 OR S2 OR S3) AND S5
S7
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                S6 NOT S4
S8
            6
                S7/2000:2003
S9
           38
                S7 NOT S8
?s (s1 or s2 or s3)
            8257 S1
            4441 S2
             779
                  S3
     S10
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?s s10 and (western? or immunoblot? or blot?)
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          124711 WESTERN?
           53434 IMMUNOBLOT?
          170904 BLOT?
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           44622 KD
           12846 DALTON?
            6642 KILODALTON?
              24 RMW
          550619 WEIGHT?
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>>>Unmatched parentheses
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               IGG?))
>>>Unrecognizable Command
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           53949 ANTISER?
           76481 IGG
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                 TGA
           39527
                 TGM
               1 MMUNE?
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                 SEROTYP?
          237721
                  IMMUNE?
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                  IMMUNOGLOB?
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                  MMUNE? OR SEROTYP? OR IMMUNE? OR IMMUNOGLOB?)
?s s13 and (avian? or fowl? or poultry? or turkey? or chick? or hen? or yolk? or igy or b
ird?)
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            5994 FOWL?
           21203 POULTRY?
           15337 TURKEY?
          127853 CHICK?
           72384 HEN?
           10851 YOLK?
             250 IGY
           34792 BIRD?
     S14
               9 S13 AND (AVIAN? OR FOWL? OR POULTRY? OR TURKEY? OR CHICK?
                  OR HEN? OR YOLK? OR IGY OR BIRD?)
?t s14/9/all
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	WEST	
	Generate Collection Print	Coupeling
L4: Entry 5 of 7	File: USPT	Apr 12, 1994

DOCUMENT-IDENTIFIER: US 5302388 A

TITLE: Control of campylobacter jejuni colonization

CLAIMS:

- 1. An anti-Campylobacter jejuni colonizing poultry feed which is useful for preventing the colonization of Campylobacter jejuni in a poultry animal, having dispersed therein, as an active ingredient, an effective amount of at least one cecal-colonizing strain of microorganism to provide anti-Campylobacter activity, wherein the strain is selected from the group consisting of Klebsiella pneumoniae strain 23 (ATCC No. 55234), Citrobacter diversus strain 22 (ATCC No. 55236), Escherichia coli (013:H.sup.-) strain 25 (ATCC No. 55235), mutants thereof which retain the ability to produce anti-Campylobacter activity, and mixtures thereof.
- 7. The composition of claim 5, wherein the cecal-colonizing strain is an active ingredient in a <u>poultry</u> feed material, said strain producing anti-<u>Campylobacter</u> metabolites to provide, upon addition to a <u>poultry</u> feed, a <u>poultry</u> feed producing an effective amount of anti-<u>Campylobacter</u> metabolites to inhibit the colonization of <u>Campylobacter</u> jejuni in a <u>poultry</u> animal.
- 11. A process for inhibiting the colonization of <u>Campylobacter jejuni in poultry</u> comprising administering an effective amount of at least one cecal-colonizing strain of microorganism, said strain producing anti-<u>Campylobacter</u> metabolites, wherein the strain is selected from the group consisting of Klebsiella pneumoniae strain 23 (ATCC No. 55234), Citrobacter diversus strain 22 (ATCC No. 55236), Escherichia coli (013:H.sup.-) strain 25 (ATCC No. 55235), mutants thereof which retain the ability to produce anti-<u>Campylobacter</u> activity, and mixtures thereof, and a carrier.
- 20. A process for inhibiting the colonization of <u>Campylobacter jejuni in poultry</u> comprising dispensing and delivering a dietary supplement comprising an effective amount of at least one cecal-colonizing strain of microorganism, said strain producing anti-<u>Campylobacter</u> metabolites, wherein the strain is selected from the group consisting of Klebsiella pneumoniae strain 23 (ATCC No. 55234), Citrobacter diversus strain 22 (ATCC No. 55236), Escherichia coli (013:H.sup.-) strain 25 (ATCC No. 55235), mutants thereof which retain the ability to produce anti-<u>Campylobacter</u> activity, and mixtures thereof.

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L8: Entry 10 of 16

File: USPT

Mar 30, 1999

DOCUMENT-IDENTIFIER: US 5888810 A

TITLE: Campylobacteri jejuni flagellin-escherichia coli LT-B fusion protein

Brief Summary Text (33):

Serum antibody response to invasive enteric pathogens is very important in protection against systemic infections. The initial immunologic response to enteric infection occurs at the level of the intestinal mucosa. Secretory immunoglobulin A (sIgA) response at the intestinal mucosa is a primary defense against enteric infections (Winsor et al. supra). Stern et al. (1990. <u>Avian</u> Dis., vol. 34, pp. 595-601) found that specific anti-C. jejuni antibodies diminish the ability of the bacterium to colonize the gut of 1-day-old <u>chicks</u> when incubated with the organism as compared with preincubation with phosphate buffered saline.

Brief Summary Text (34):

The flagella of C. jejuni are essential in the colonization of the intestine. Nonflagellated organisms are quickly cleared from the intestine. <u>Chicken polyclonal</u> antiflagellin antibodies as well as <u>monoclonal</u> antiflagellin antibodies have been found to prevent C. jejuni from colonizing the <u>chickens</u> or to increase the dose of bacteria required to colonize the <u>chickens</u> (Carr, unpublished). Flagellar antigens are therefore potential candidates for vaccines as well as suitable antigens for diagnostic purposes, since the flagellin protein is immunodominant during human infections.

Detailed Description Text (9):

The LT-B/fla fusion gene was under constitutive expression in X6097. The fusion protein was detected at several growth times. The best recovery, i.e. the greatest yield of the fusion protein relative to the total protein, was when cell density corresponding to OD.sub.600 of about 0.8 was reached. The fusion protein was detected by Coomassie staining, and Western blot analyses using <u>chicken</u> anti-flagellin serum (FIG. 2), rabbit anti-C. <u>jejuni</u> serum, affinity purified rabbit anti-C. <u>jejuni</u> flagellin antibodies and rabbit anti-LT serum (FIG. 2). The fusion protein was not recognized by a <u>monoclonal</u> antibody directed against the 63 kd flagellin protein, presumably because the <u>monoclonal</u> antibody is directed against an epitope not present in our fusion protein since only 46% of the flaA gene is expressed. The fusion protein has a MW of 43 kd (16 for LT-B and 27 kd for the U band). The protein could not be detected from the pBEB transformed X6097 control. The LT-B/fla, fusin gene DNA sequence is presented in FIG. 3.

<u>Detailed Description Text</u> (28):

Western blot analyses were performed as described by Towbin et al (1979. PNAS, vol. 76, pp. 4350-4354). Blots were treated with a 1:200 dilution of <u>chicken</u> anti-*C.* <u>jejuni</u> serum or rabbit anti-LT before adding the secondary antibody (goat-anti rabbit <u>IgG</u> alkaline phosphatase conjugate, Bio-Rad), and developed with the substrate solution (5-bromo-4-chloro-3-indolyl phosphate/nitro blue tetrazolium) as described by Sambrook et al. (1989. In Molecular Cloning, A Laboratory Manual, Cold Spring Harbor, N.Y.: Cold Spring Harbor

Laboratory). Results are shown in FIG. 2.

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S2/2000:2003
 S3
           295
 S4
           78 S2 NOT S3
 ?t s4/3, kwic/42
 4/3, KWIC/42
                 (Item 17 from file: 349)
DIALOG(R) File 349: PCT FULLTEXT
 (c) 2003 WIPO/Univentio. All rts. reserv.
00452378
            **Image available**
A PORIN GENE FROM CAMPYLOBACTER JEJUNI, RELATED PRODUCTS AND USES THEREOF
GENE DE PORINE EXTRAIT DE CAMPYLOBACTER JEJUNI, PRODUITS APPARENTES ET
    LEURS UTILISATIONS
Patent Applicant/Assignee:
  HER MAJESTY IN RIGHT OF CANADA as represented by THE MINISTER OF HEALTH
    AND WELFARE CANADA,
  JOHNSON Wendy M,
  BACON David J,
  RODGERS Frank,
  BOLLA Jean-Michel,
Inventor(s):
  JOHNSON Wendy M,
  BACON David J.
  RODGERS Frank.
  BOLLA Jean-Michel,
Patent and Priority Information (Country, Number, Date):
  Patent:
                      WO 98CA272 19980325
                        WO 9842842 A1 19981001
  Application:
                                             (PCT/WO CA9800272)
  Priority Application: US 9741200 19970325
Designated States: AL AM AT AU AZ BA BB BG BR BY CA CH CN CU CZ DE DK EE ES
  FI GB GE GH GM GW HU ID IL IS JP KE KG KP KR KZ LC LK LR LS LT LU LV MD
  MG MK MN MW MX NO NZ PL PT RO RU SD SE SG SI SK SL TJ TM TR TT UA UG US
  UZ VN YU ZW GH GM KE LS MW SD SZ UG ZW AM AZ BY KG KZ MD RU TJ TM AT BE
  CH DE DK ES FI FR GB GR IE IT LU MC NL PT SE BF BJ CF CG CI CM GA GN ML
  MR NE SN TD TG
Publication Language: English
Fulltext Word Count: 25116
A PORIN GENE FROM CAMPYLOBACTER JEJUNI, RELATED PRODUCTS AND USES THEREOF
GENE DE PORINE EXTRAIT DE CAMPYLOBACTER JEJUNI, PRODUITS APPARENTES ET
    LEURS UTILISATIONS
Fulltext Availability:
  Detailed Description
  Claims
English Abstract
  The invention relates to a porin gene from Campylobacter jejuni [SEQ ID
  NO:3]. The gene has been designated porA and is 1275 bp...
French Abstract
  L'invention concerne un gene de porine extrait de Campylobacter jejuni
  (SEQ ID NO:3). Ce gene, denomme porA, a une longueur de 1275 bp...
Detailed Description
  A PORIN GENE FROM Campylobacter jejuni,
  RELATED PRODUCTS AND USES THEREOF
  TECHNICAL FIELD
  This invention relates to a porin gene from
  5 Campylobacter jejuni, to related products and to the uses
  BACKGROUND ART
  In the following discussion, the...
...41). Active surveys
 conducted in the United States have estimated the number
 of cases of campylobacteriosis to be 2.5 million per year,
 making it a multi-million dollar disease (39...
...by C. jejuni can range from watery to bloody
```

diarrhea (28, 39). In most cases campylobacteriosis is a

self-limiting disease but in the more severe cases,

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File 155:MEDLINE(R) 1966-2003/Aug W3
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*File 155: Medline has been reloaded and accession numbers have changed. Please see HELP NEWS 155.

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Set Items Description
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?e campylobacter

Ref	Items	RT Index-term
E1	1	CAMPYLOBACHTER
E2	1	CAMPYLOBACTE
E3	8257	9 *CAMPYLOBACTER
E4	54	CAMPYLOBACTERANALYSISAN
E5	21	CAMPYLOBACTERCHEMISTRYCH
E6	529	CAMPYLOBACTERCLASSIFICATIONCL
E7	15	CAMPYLOBACTERCYTOLOGYCY
E8	398	CAMPYLOBACTERDRUG EFFECTSDE
E9	125	CAMPYLOBACTERENZYMOLOGYEN
E10	317	CAMPYLOBACTERGENETICSGE
E11	270	CAMPYLOBACTERGROWTH AND DEVELOPMENTGD
E12	295	CAMPYLOBACTERIMMUNOLOGYIM

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E15
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E27	148		CAMPYLOBACTER COLIISOLATION AND PURIFICATIO	
E28	17		CAMPYLOBACTER COLIMETABOLISMME	
E29	24		CAMPYLOBACTER COLIPATHOGENICITYPY	
E30	16		CAMPYLOBACTER COLIPHYSIOLOGYPH	
E31.	8		CAMPYLOBACTER COLIULTRASTRUCTUREUL	
E32	1772	5	CAMPYLOBACTER FETUS	
E33	35		CAMPYLOBACTER FETUSANALYSISAN	
E34	14		CAMPYLOBACTER FETUSCHEMISTRYCH	
E35	192		CAMPYLOBACTER FETUSCLASSIFICATIONCL	
E36	14		CAMPYLOBACTER FETUSCYTOLOGYCY	

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E39	105		CAMPYLOBACTER	FETUS	GENETICSGE
E40	157		CAMPYLOBACTER	FETUS	GROWTH AND DEVELOPMENT -
E41	241		CAMPYLOBACTER	FETUS	IMMUNOLOGYIM
E42	781		CAMPYLOBACTER	FETUS	ISOLATION AND PURIFICATI
E43	80		CAMPYLOBACTER	FETUS	METABOLISMME

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108
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             CAMPYLOBACTER INFECTIONS -- CEREBROSPINAL FLUID
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E47
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E48
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                          affecting
     study of
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                                       the sensitivity of the passive
haemagglutination method
                           for serotyping Campylobacter jejuni and
Campylobacter coli and recommendations for a more rapid procedure.
 Fricker C R; Alemohammad M M; Park R W
 Canadian journal of microbiology (CANADA) Jan 1987, 33 (1) p33-9,
Document type: Journal Article
 Languages: ENGLISH
 Main Citation Owner: NLM
 Record type: Completed
 Subfile:
            INDEX MEDICUS
 Factors affecting the sensitivity of the passive haemagglutination method
for serotyping campylobacters have been studied. The concentration of red
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2 CAMPYLOBACTERIUM

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blood cells during the haemagglutination stage of the procedure markedly
 affected the titer obtained. An increase in concentration of red blood
cells resulted in a lower titer, with titers being inversely proportional
to red blood cell concentration. No differences in titer were observed when
erythrocytes were sensitized at a range of pH values between pH 5.0 and pH
8.0. The time required for antigen extraction and for red blood cell
sensitization was shown to be 15 min each, thus resulting in a reduction in
      time required
                              serotyping . Furthermore, use of
                      for
erythrocytes enabled the haemagglutination reactions to be read after
incubation for only 1 h. Combining these procedures with a rapid slide
haemagglutination test enables a single worker to serotype over 100 C.
jejuni and C. coli isolates within 1 working day.
  Tags: Animal; Comparative Study; Human; Support, Non-U.S. Gov't
  Descriptors: Campylobacter -- classification -- CL; * Campylobacter fetus
--classification--CL;
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                                                                Antigens,
            Chickens--blood--BL; Hydrogen-Ion Concentration; Serotyping
--methods--MT; Sheep--blood--BL; Time Factors; Turkeys--blood--BL
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*File 155: Medline has been reloaded and accession numbers have
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  File 340:CLAIMS(R)/US Patent 1950-03/Aug 14
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  File 444: New England Journal of Med. 1985-2003/Aug W3
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       16:Gale Group PROMT(R) 1990-2003/Aug 15
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*File 16: Alert feature enhanced for multiple files, duplicate
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S2

373 S1 AND CAMPYLOBAC?

antibiotic intervention with macrolids...

...34). Johnson

fragment thereof.

5 and Lior (19) originally reported that 410@o of 718 isolates of **Campylobacter** sp. screened for the production of CLDT were positive; however, isolates screened for the cdt13...said target.

The invention also relates to a method of detecting 25 the presence of **Campylobacter** jejuni infection, characterized by the steps of: a) contacting a sample obtained from a patient...

...a time sufficient to allow formation of a complex between said protein and any anti- Campylobacter jejuni antibodies present in said sample; and b) detecting the presence of, and optionally the...

...comprises an isolated expression vector, characterized by a region encoding a -5 porA protein of Campylobacter jejuni, or an antigenic

Included within the invention is a method of inducing... ... aspect of the invention is a method of producing antibodies for testing for infection by Campylobacter jejuni, characterized in that a protein having an amino acid sequence of SEQ ID NO...h; Figure 8 shows western blot analysis of the isolated cytotoxic porin-LPS complex from Campylobacter sp using 40 Ag of crude, concentrated filtrate and homologous rabbit antiserum. Lanes 1 and...present invention is based on the identification 5 of a porin-lipopolysaccharide (LPS) complex from Campylobacter jejuni that is an endotoxin and that is fairly well conserved amongst strains of the organism, but not widely found in other Campylobacter species. The complex has been isolated and a corresponding porin gene, 10 designated 11porA,11...subsequently stored at -700C in try-ptic soy broth

and related organisms were maintained at -800C in glycerol-peptone water as part of...ompC of S. typhi (29), a 45i similarity and 200i identity 25 was found.

Screening Campylobacter sp. for porA and cytotoxin production.

containing 5% sheep blood. Strains of Campylobacter sp.

Results of screening C. jejuni for phenotypic ...gene are summarized 30 in Table 2. It was found that all 32 strains of Campylobacter sp. and related organisms produced a cytotoxic component when the filtrate from the biphasic -44...

...of C. jejuni, especially Lior serotype 82, but was not conserved between related species of Campylobacter.

DISCUSSION

The 1275 bp ORF had a '@.guanosine+cytosine content of 36.8 mol % (Fig...contained at least

part, if not all of the intact gene while the other Campylobacter sp. and related organism were PCR negative.

Previous reports indicated that only 60'i of...
...are valuable, and provide a new and
efficient method to identify C. jejuni from other

Campylobacter sp. The potential for the development of a recombinant vaccine using the porin protein is...on immuno-blot analysis. Western blots of crude concentrated filtrates from various cytotoxic strains of Campylobacter species showed the presence of a protein with a molecular mass similar to that of...comparable molecular weight was also present in crude concentrated filtrates from other cytotoxic strains of Campylobacter sp., indicating that the release of the porin-LPS complex 25 was not unique to...been cloned and sequenced, a fuller understanding of the role of the porin in clinical campylobacteriosis will be forthcoming. Such evaluations may suggest potential roles for the porin-LPS complex as...COUNTRY: France (F) POSTAL CODE (ZIP): 13009

- (ii) TITLE OF INVENTION: A PORIN GENE FROM CAMPYLOBACTER JEJUNI, RELATED PRODUCTS AND USES THEREOF
- (iii) NUMBER OF SEQUENCES: 31
- -70 (iv) COMPUTER READABLE...unknown
- (ii) MOLECULE TYPE: protein
- (v) FRAGMENT TYPE: N-terminal
- (vi) ORIGINAL SOURCE.
- (A) ORGANISM: Campylobacter jejuni
- (B) STRAIN: 2483
- (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 2.

Met Lys Leu Val...STRANDEDNESS: double

- (D) TOPOLOGY: linear
- (ii) MOLECULE TYPE: DNA (genomic)
- (vi) ORIGINAL SOURCE.
- (A) ORGANISM: Campylobacter jejuni
- (B) STRAIN: 2483
- (xi) SEQUENCE DESCRIPTION: SEQ ID NO: 3.

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Dis. 1997; 176 (Suppl 2): S125

Claim

- ... purif ied nucleic acid, characterized in that said nucleic acid encodes a porA protein of Campylobacter jejuni, or an antigenic fragment thereof.
 - 2 A nucleic acid according to claim 1, characterized...
- ...acid according to claim 1, characterized in that it is derived from strain 2483 of **Campylobacter** 10 jejuni (ATCC Accession No.
 - 4 A nucleic acid according to claim 1, characterized in...
- ...probe

to bind specifically to said target.

- 59 A method of detecting the presence of **Campylobacter** jejuni infection, characterized by the steps of:
 a) contacting a sample obtained from a patient...
- ...time sufficient to
 - 10 allow formation of a complex between said protein and any anti- Campylobacter jejuni antibodies present in said sample; and
 - b) detecting the presence of, and optionally the...
- ...said complex formed during step (a).
 15 10. A method of detecting the presence of Campylobacter jejuni in a patient, characterized by obtaining from said patient a sample suspected of containing Campylobacter jejuni, and detecting whether the characteristic nucleic acid of claim 1, claim 2, claim 3...

...103

- 14 An isolated expression vector, characterized by a region encoding a porA protein of **Campylobacter** jejuni, or an antigenic ...said complex.
- 19 A vaccine comprising an immunogenically effective amount of the porA antigen of **Campylobacter** ' ' . or 3e3un7

antigenic fragment thereof and a pharmaceutically acceptable carrier.

20 20. A vaccine, characterized...

...human or animal body.

22 A method of producing antibodies for testing for infection by **Campylobacter** jejuni, characterized in that a 5 protein having an amino acid sequence of SEQ ID... ?t s4/9/62 64 68

4/9/62 (Item 3 from file: 5)
DIALOG(R)File 5:Biosis Previews(R)
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10109393 BIOSIS NO.: 199698564311

Identification and characterization of an immunogenic outer membrane protein of Campylobacter jejuni.

AUTHOR: Burnens Andre; Stucki Urs; Nicolet Jacques; Frey Joachim(a) AUTHOR ADDRESS: (a) Inst. Veterinary Bacteriol., Univ. Berne,

Langgassstrasse 122, CH-3012 Berne**Switzerland

JOURNAL: Journal of Clinical Microbiology 33 (11):p2826-2832 1995

ISSN: 0095-1137

DOCUMENT TYPE: Article RECORD TYPE: Abstract LANGUAGE: English

ABSTRACT: We cloned and expressed in Escherichia coli a gene encoding an 18- kDa outer membrane protein (Omp18) from Campylobacter jejuni ATCC 29428. The nucleotide sequence of the gene encoding Omp18 was determined, and an open reading frame of 165 amino acids was revealed. The amino acid sequence had the typical features of a leader sequence and a signal peptidase II cleavage site at the N-terminal part of Omp18. Moreover, the sequence had a high degree of similarity to the peptidoglycan-associated outer membrane lipoprotein P6 of Haemophilus influenzae and the peptidoglycan-associated lipoprotein PAL of E. coli. Southern blot analysis in which the cloned gene was used as a probe revealed genes similar to that encoding Omp18 in all species of the thermophilic group of campylobacters as well as Campylobacter sputorum. All campylobacters tested expressed a protein with a molecular mass identical to that of Omp18. The protein reacted immunologically with antibodies directed against Omp18 from C. jejuni. PCR amplification of the gene encoding Omp18 with specific primers and subsequent restriction enzyme analysis of the amplified DNA fragments showed that the gene for Omp18 is highly conserved in C. jejuni strains isolated from humans, dogs, cats, calves, and chickens but is different in other Campylohacter species. In order to obtain pure recombinant Omp18 protein for serological assays, the cloned gene for Omp18 was genetically modified by replacing the signal sequence with a DNA segment encoding six adjacent histidine residues. Expression of this construct in E. coli allowed showed no reaction with this antigen. Omp18, which is an outer membrane protein belonging to the family of PALs is well conserved in C. jejuni and is highly immunogenic. It is therefore a good candidate as an antigen for the serological diagnosis of past C. jejuni infections.

REGISTRY NUMBERS: 170613-04-4: GENBANK-X83374 DESCRIPTORS:

MAJOR CONCEPTS: Biochemistry and Molecular Biophysics; Blood and Lymphatics (Transport and Circulation); Immune System (Chemical Coordination and Homeostasis); Infection; Membranes (Cell Biology); Pathology; Veterinary Medicine (Medical Sciences)

BIOSYSTEMATIC NAMES: Aerobic Helical or Vibrioid Gram-Negatives-Eubacteria, Bacteria; Bovidae--Artiodactyla, Mammalia, Vertebrata,
Chordata, Animalia; Canidae--Carnivora, Mammalia, Vertebrata, Chordata,
Animalia; Felidae--Carnivora, Mammalia, Vertebrata, Chordata, Animalia;
Galliformes--Aves, Vertebrata, Chordata, Animalia; Hominidae--Primates,
Mammalia, Vertebrata, Chordata, Animalia; Pasteurellaceae--Eubacteria,
Bacteria; Trogoniformes--Aves, Vertebrata, Chordata, Animalia

ORGANISMS: aerobic helical or vibrioid gram-negative bacteria (Aerobic Helical or Vibrioid Gram-Negatives); calf (Bovidae); cat (Felidae); chicken (Galliformes); dog (Canidae); human (Hominidae); Campylobacter jejuni (Aerobic Helical or Vibrioid Gram-Negatives); Campylobacter sputorum (Aerobic Helical or Vibrioid Gram-Negatives); Haemophilus influenzae (Pasteurellaceae); Trogoniformes (Trogoniformes

BIOSYSTEMATIC CLASSIFICATION (SUPER TAXA): animals; artiodactyls; bacteria; birds; carnivores; chordates; eubacteria; humans; mammals; microorganisms; nonhuman mammals; nonhuman vertebrates; primates; vertebrates

CHEMICALS & BIOCHEMICALS: GENBANK-X83374

MOLECULAR SEQUENCE DATABANK NUMBER: molecular sequence data; nucleotide sequence; EMBL-X83374; GENBANK-X83374

MISCELLANEOUS TERMS: INFECTION; OPEN READING FRAME; OUTER MEMBRANE PROTEIN 18 GENE; PEPTIDOGLYCAN-ASSOCIATED OUTER MEMBRANE LIPOPROTEIN P6 SIMILARITY; SEROLOGICAL DIAGNOSIS CONCEPT CODES:

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$2.54 Estimated cost File349
       $0.30
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          $3.50 2 Type(s) in Format 9
       $3.50 2 Types
$3.80 Estimated cost File5
       $0.04 0.009 DialUnits File35
$0.04 Estimated cost File35
       $0.05 0.018 DialUnits File10
          $1.35 1 Type(s) in Format 9
       $1.35 1 Types
$1.40 Estimated cost File10
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       $0.10
             0.009 DialUnits File347
$0.10 Estimated cost File347
       $0.05 0.009 DialUnits File160
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       $0.02 0.009 DialUnits File143
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       $0.03 0.009 DialUnits File370
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      OneSearch, 21 files, 0.430 DialUnits FileOS
$0.22 TELNET
$8.85 Estimated cost this search
$8.85 Estimated total session cost 0.430 DialUnits
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Status: Signed Off. (1 minutes)

10973131 97325879 PMID: 9182891

Oral administration of antibodies as prophylaxis and therapy in

Campylobacter jejuni-infected chickens.

Tsubokura K; Berndtson E; Bogstedt A; Kaijser B; Kim M; Ozeki M; Hammarstrom L

Department of Clinical Immunology, Huddinge Hospital, Sweden.

Clinical and experimental immunology (ENGLAND) Jun 1997, 108 (3) p451-5, ISSN 0009-9104 Journal Code: 0057202

Document type: Journal Article

Languages: ENGLISH

Main Citation Owner: NLM Record type: Completed Subfile: INDEX MEDICUS

Passive immunity against gastrointestinal infections has recently been successfully applied as prophylaxis and therapy in patients in a variety of virally and bacterially induced infections. Campylobacter jejuni is frequently associated with acute diarrhoea in humans, and several species of animals have been shown to transmit the disease, although birds have been implicated as the main source of infection. We used bovine and chicken immunoglobulin preparations from the milk and eggs, respectively, of immunized animals for prophylactic and therapeutic treatment of chickens infected with C. jejuni. A marked prophylactic effect (a >99% decrease in the number of bacteria) was noted using either antibody preparation, whereas the therapeutic efficacy, i.e. when antibodies were given after the infection was established, was distinctly lower (80-95%) as judged by faecal bacterial counts. These observations may serve as a starting point for experiments aimed at elimination of the infection in an industrial or farm setting. It may also encourage future attempts to treat, prophylactically or therapeutically, patients with Campylobacter -induced diarrhoea.

Tags: Animal; Female; Support, Non-U.S. Gov't

Descriptors: Antibodies, Bacterial--therapeutic use--TU; * Campylobacter Infections--prevention and control--PC; * Campylobacter --immunology--IM; Administration, Oral; Campylobacter Infections--therapy

--TH; Cattle; Chickens; Immunization, Passive CAS Registry No.: 0 (Antibodies, Bacterial)

Record Date Created: 19970626 Record Date Completed: 19970626 11654827 99089498 PMID: 9874102

The specificity of antibody in chickens immunised to reduce intestinal colonisation with Campylobacter jejuni.

Widders P R; Thomas L M; Long K A; Tokhi M A; Panaccio M; Apos E
Australian Quarantine and Inspection Service, Mascot, NS
phillip.widders@dpie.qov.au

Veterinary microbiology (NETHERLANDS) Nov 1998, 64 (1) p39-50,

ISSN 0378-1135 Journal Code: 7705469

Document type: Journal Article

Languages: ENGLISH
Main Citation Owner: NLM
Record type: Completed
Subfile: INDEX MEDICUS

Poultry consumption has been identified as a major risk factor for human infection with Campylobacter jejuni in developed countries. C. jejuni is present in the gastrointestinal tract of broiler chickens at the time of slaughter, and faecal contamination of carcases during processing results in significant campylobacter loads on carcases. One approach to reducing the level of carcase contamination with C. jejuni is to control campylobacter infection in broiler chickens. To this end, the study described here investigated the specificity of antibody in serum and intestinal secretions of chickens that had been immunised with antigens and then challenged with viable bacteria. The campylobacter immunodominant antigens in the serum of **birds** that showed a 2-log reduction in caecal colonisation with C. jejuni included flagellin protein Kd) and three additional antigens of 67, 73.5 and 77.5 Kd . Only flagellin and the 67 Kd antigen were recognised by IgG antibody in gastrointestinal secretions of the same birds . Antibody from chickens immunised with purified native flagellin protein recognised flagellin protein and the 67 Kd antigen in Western blots probed with serum, but only the flagellin proteins (61-63 Kd) in Westerns probed with gastrointestinal secretions. Analysis of the specificity of the response to flagellin protein using recombinant clones that expressed regions of the flagellin gene suggests that epitopes in each region of the flagellin protein were immunogenic. Of the immunodominant antigens, only flagellin appeared to be surface-exposed on viable C. jejuni, although conformational epitopes of flagellin appeared to be sensitive to the method of antigen purification. The results of this study suggest that flagellin and possibly the 67 Kd antigen may be valuable for immunological control of intestinal infection with C. jejuni in chickens , but that further work is required to purify these as vaccine candidates by using methods that preserve conformational epitopes.

Tags: Animal; Support, Non-U.S. Gov't

Descriptors: Antibodies , Bacterial--immunology--IM; * Antibody Specificity; * Campylobacter Infections--veterinary--VE; * Campylobacter

11829439 99269532 PMID: 10337238

Biotin-streptavidin enzyme-linked immunosorbent assay for the detection of antibodies to Campylobacter jejuni and C. coli in chickens.

Haas B; Hinz K H; Glunder G

Clinic for Poultry, Hanover School of Veterinary Medicine, Germany.

Zentralblatt fur Veterinarmedizin. Reihe B. Journal of veterinary medicine. Series B (GERMANY) Apr 1999, 46 (3) p163-71, ISSN 0514-7166

Journal Code: 0331325

Document type: Journal Article

Languages: ENGLISH

Main Citation Owner: NLM Record type: Completed Subfile: INDEX MEDICUS

immunosorbent assay (ELISA) was developed in a enzyme-linked homologous system with bacterial ultrasonic-treated proteins as the antigen and antisera from chickens infected orally and subcutaneously with the strain Campylobacter jejuni serovar 6 (CJ 6). The cut-off level was determined using antisera from non-infected specific-pathogen-free chickens up to the age of 10 weeks. The suitability of the ELISA system was verified antisera taken from chickens orally infected at the age of 4 weeks with CJ 1, 6, 28 or 36 or with Campylobacter coli serovar 28 (CC development of antibodies was monitored up to 6 weeks post-infection (p.i.). Sera from chickens infected with CJ 1, 6, 36 or CC 28 contained specific antibodies to **Campylobacter** , whereas in those infected with CJ 28 no specific antibodies were found. Distinct cross-reactions were observed between CJ 6, 28 and CC 28 antigens and their antisera 6 weeks p.i., while poor cross-reactions were found with antisera to CJ 1 and 28. Antibodies to strains of all heterologous serovars were successfully detected with an antigen pool comprised of CJ 1, 6 and 36 antigens. In 11 out of the 12 field sera obtained from 5- and 9-week-old broiler chickens suffering from campylobacteriosis , high specific antibody titres to Campylobacter jejuni were found. Tags: Animal

10700234 '97049491 PMID: 8894221

Immunisation of chickens to reduce intestinal colonisation with Campylobacter jejuni.

Widders P R; Perry R; Muir W I; Husband A J; Long K A

Department of Agriculture, Energy and Minerals, Victorian Institute of Animal Science, Attwood, Australia.

British poultry science (ENGLAND) 07-1668 Journal Code: 15740290R Sep 1996, 37 (4) p765-78, ISSN

Document type: Journal Article

Languages: ENGLISH

Main Citation Owner: NLM Record type: Completed INDEX MEDICUS

1. Systemic and intestinal antibody titres were measured in chickens following subcutaneous, intraperitoneal (i.p.), oral (p.o.) and combined administration of antigen, in soluble, emulsified or microparticulate form. Antigens tested included keyhole limpet haemocyanin killed Campylobacter jejuni whole cells and purified campylobacter flagellin protein. 2. The effect of immunisation with purified flagellin protein or with killed C. jejuni whole cells in reducing intestinal colonisation was assessed. The ability of newlyhatched chicks to respond to immunisation was limited, possibly because of the immaturity of the immune system rather than maternal suppression of an immune response. Only 5 to 13 birds that were first immunised when 1-d-old with KLH showed a systemic response, even after 4 immunisations, whereas 10 of 11 birds that were first immunised at 24 d-old responded systemically. 3. In an immunisation and challenge experiment, birds that were immunised twice intraperitoneally, at 16 and 29 d-old, with killed C. jejuni whole cells, had fewer C. jejuni, in the caecal contents than unimmunised control birds. This reduction in intestinal colonisation, to less than 2% of bacterial numbers in control birds, was associated with an increase in specific IgG in intestinal secretions. There was no significant increase in specific IgA or IgM in intestinal secretions following immunisation and challenge. 4. These results indicate that immunisation can reduce the level of intestinal infection with C. jejuni. The protection may be enhanced by developing improved methods of immunisation that stimulate production of increased titres of specific antibody in intestinal secretions, particularly specific IgA antibody.

Tags: Animal; Female; Support, Non-U.S. Gov't

Descriptors: Antibodies, Bacterial--metabolism--ME; * Campylobacter Infections--veterinary--VE; * Campylobacter jejuni --isolation and purification--IP; *Chickens--immunology--IM; *Intestines--microbiology--MI; *Poultry Diseases--immunology--IM; *Poultry Diseases --prevention and control--PC; *Vaccines, Inactivated--pharmacology--PD; Antibodies, Bacterial--immunology--IM; Antigens--immunology--IM; Antigens--metabolism --ME; Antigens, Bacterial--immunology--IM; Antigens, Bacterial--metabolism Infections -- immunology -- IM; Campylobacter Campylobacter Infections--prevention and control--PC; Campylobacter jejuni--immunology Feces--microbiology--MI; Flagellin--immunology--IM; --pharmacology--PD; Hemocyanin--immunology--IM; Hemocyanin--pharmacology --PD; Immunoglobulin A--immunology--IM; Immunoglobulin A--metabolism--ME; Immunoglobulin G--immunology--IM; Immunoglobulin G--metabolism--ME; Intestines--drug effects--DE; Time Factors; Vaccines, Inactivated --immunology--IM

Registry No.: 0 (Antibodies, Bacterial); 0 (Antigens); 0 (Antigens, Bacterial); 0 (Immunoglobulin A); 0 (Immunoglobulin G); 0 (Vaccines, Inactivated); 12777-81-0 (Flagellin); 9013-72-3 (Hemocyanin) Record Date Created: 19970206

Record Date Completed: 19970206

10291007 96092877 PMID: 8590089

In ovo oral vaccination with Campylobacter jejuni establishes early development of intestinal immunity in chickens.

Noor S M; Husband A J; Widders P R

Department of Veterinary Pathology, University of Sydney, New South Wales, Australia.

British poultry science (ENGLAND) Sep 1995, 36 (4) p563-73, ISSN 007-1668 Journal Code: 15740290R 0007-1668

Document type: Journal Article

Languages: ENGLISH

Main Citation Owner: NLM Record type: Completed Subfile: INDEX MEDICUS

Chick embryos were orally immunised at day 16 of incubation by injection of heat-killed **Campylobacter** jejuni organisms into the amniotic fluid. The response to vaccination was observed at 5 d after hatching or, in some birds which received a postnatal oral booster vaccination, at 7 d after hatching, and the response was observed at 14 d of age. 2. The titres of antibody in serum, bile and intestinal scrapings, the distribution of immunoglobulin-containing cells in the spleen, duodenum and ileum and the expression on peripheral blood leukocytes (PBL) of the T cell surface markers CD3, CD4 and CD8 were determined. 3. Whereas low titres of anti-flagellin antibody were detected in serum, bile and intestinal scrapings of unimmunised birds, high titres were observed in immunised birds. 4. An increase in antibody of all isotypes was detectable in serum but the elevation in IgA antibody in intestinal scrapings and bile was particularly striking. This response was reflected in a dramatic in immunoglobulin-containing cells, detected by fluorescent histology, particularly those associated with IgA and IgM isotypes in the spleen and intestine of immunised birds. 5. Secondary oral boosting after hatching resulted in a depression in serum anti-flagellin antibody in immunised birds compared to pre-boosting titres (although still significantly higher than in non-immunised controls) but an increase in IgA intestinal scrapings in and bile. The number immunoglobulin-containing cells was also increased after boosting. 6. Neither immunisation regimen caused a significant change in the numbers of circulating CD3, CD4 or CD8 T cells. 7. These results indicate that in ovo oral immunisation with C. jejuni antigens stimulates the precocious development of immunity in chicks.

Tags: Animal; Support, Non-U.S. Gov't

Descriptors: Campylobacter Infections -- veterinary -- VE; * Campylobacter jejuni--immunology--IM; *Chickens--immunology--IM; *Intestines--immunology *Poultry Diseases--immunology--IM; *Vaccination--veterinary--VE; Administration, Amnion--immunology--IM; Oral: Antibodies, --analysis--AN; Antibodies, Bacterial--blood--BL; Antigens, CD3--analysis Antigens, CD4--analysis--AN; Antigens, CD8--analysis--AN; Bile --immunology--IM; Campylobacter Infections--immunology--IM; Campylobact Infections -- prevention and control -- PC; Chickens -- metabolism -- ME; Chickens--microbiology--MI; Immunity, Mucosal; Immunization, Secondary-veterinary--VE; Immunoglobulins--blood --veterinary--VE; --BL; Intestines--microbiology--MI; Lymphocytes--cytology--CY; Lymphocytes --immunology--IM; Poultry Diseases--prevention and control--PC; Vaccination --methods--MT; Weight Gain--physiology--PH CAS Registry No.: 0 (Antibodies, Bacterial); 0 (Antigens, CD3); 0

(Antigens, CD4); 0 (Antigens, CD8); 0 (Immunoglobulins)

Record Date Created: 19960325 Record Date Completed: 19960325 08383145 95071121 PMID: 7526839

Isotype, specificity, and kinetics of systemic and mucosal antibodies to Campylobacter jejuni antigens, including flagellin, during experimental oral infections of chickens.

Cawthraw S; Ayling R; Nuijten P; Wassenaar T; Newell D G

Central Veterinary Laboratory (Weybridge), New Haw, Addlestone, Surrey, United Kingdom.

Avian diseases (UNITED STATES) Apr-Jun 1994, 38 (2) p341-9, ISSN 0005-2086 Journal Code: 0370617

Document type: Journal Article

Languages: ENGLISH
Main Citation Owner: NLM
Record type: Completed
Subfile: INDEX MEDICUS

The immune response of chickens to Campylobacter jejuni infection was studied as a step in the search for vaccine candidates. One-day-old chicks orally challenged with C. jejuni strain 81116 showed significant increases in specific IgG, IgA, and IgM circulating antibodies, as detected by enzyme-linked immunosorbent assay (ELISA). These levels peaked at 9, 5, and 7 weeks postinfection, respectively. Maternal IgG antibodies were also detected over the first 2 weeks. Specific mucosal IgG and IgA antibody levels also increased significantly. All of the birds demonstrated a major response to the 62-kDa flagellin protein by Western blotting techniques. The immunodominance of flagellin was confirmed by ELISA using an antigen preparation from an aflagellate mutant. When overlapping recombinant polypeptide fragments of flagellin were used, epitopes detected by chicken antibodies were observed in region IV, between residues 95-340 of the protein. Thus flagellin may be suitable candidate for a vaccine, although its role in protection must first be established.

Tags: Animal; Support, Non-U.S. Gov't

Descriptors: Antibodies , Bacterial--biosynthesis--BI; *Antigens, Bacterial--immunology--IM; * Campylobacter Infections--immunology--IM; * Campylobacter jejuni--immunology--IM; *Flagellin--immunology--IM; Immunoglobulin G--biosynthesis--BI; * Immunoglobulin Isotypes --biosynthesis--BI; *Intestinal Mucosa--immunology--IM; Antibodies , Bacterial--blood--BL; Antibodies , Bacterial -- classification -- CL; Antibody Specificity; Blotting , Western ; Chickens ; Enzyme-Linked Immunosorbent Assay; Epitopes--analysis--AN; Immunoglobulin G--blood--BL; Immunoglobulin G--classification--CL; Immunoglobulin --classification--CL CAS Registry No.: 0 (Antibodies, Bacterial); 0

CAS Registry No.: 0 (Antibodies, Bacterial); 0 (Antigens, Bacterial); 0 (Epitopes); 0 (Immunoglobulin G); 0 (Immunoglobulin Isotypes); 12777-81-0 (Flagellin)

Record Date Created: 19941129
Record Date Completed: 19941129

11197973 98074583 PMID: 9413103

Campylobacter jejuni in broiler chickens: colonization and humoral immunity following oral vaccination and experimental infection.

Rice B E; Rollins D M; Mallinson E T; Carr L; Joseph S W

Enteric Diseases Program, Naval Medical Research Institute, Bethesda, MD 20889-5607, USA.

Vaccine (ENGLAND) Dec 1997, 15 (17-18) p1922-32, ISSN 0264-410X

Journal Code: 8406899

Document type: Journal Article

Languages: ENGLISH

Main Citation Owner: NLM Record type: Completed Subfile: INDEX MEDICUS

A formalin inactivated, Campylobacter jejuni whole cell vaccine, either with or without Escherichia coli heat labile toxin (LT) as a mucosal adjuvant, was administered orally to broiler chickens. Three vaccine trials were performed, differing in the number of vaccinations, and time of administration, as well as the inclusion and dose of LT. The overall reductions of C. jejuni colonization in the vaccinated chickens ranged from 16 to 93% compared with non-vaccinated controls. Enhanced levels of anti-C. jejuni secretory IgA antibodies were demonstrated in vaccinated chickens. Vaccination also appeared to induce an anamnestic response to C. jejuni antigens in the 14-33 kDa range, as demonstrated by Western immunoblots. Interestingly, the inclusion of LT in the vaccine regimen did not appear to boost the immunogenicity of the vaccine. These results are encouraging and suggest that future development of successful oral vaccines for the control of enteropathogenic Campylobacter in poultry is feasible.

Tags: Animal; Support, Non-U.S. Gov't; Support, U.S. Gov't, Non-P.H.S. Descriptors: Bacterial Vaccines--therapeutic use--TU; * Campylobacter Infections--veterinary--VE; * Campylobacter jejuni--immunology--IM; *Poultry Diseases--immunology--IM; *Poultry Diseases--prevention and control--PC; Administration, Oral; Antibody Formation--immunology--IM; Campylobacter Infections--immunology--IM; Campylobacter Infections--prevention and control--PC; Poultry Diseases--metabolism--ME; Vaccines, Inactivated--therapeutic use--TU

CAS Registry No.: 0 (Bacterial Vaccines); 0 (Vaccines, Inactivated)

Record Date Created: 19980223
Record Date Completed: 19980223

10257830 96059438 PMID: 7483909

Development of humoral precipitating antibodies to Campylobacter spp. in chickens]

Entwicklung humoraler prazipitierender Antikorper gegen Campylobacter spp. beim Huhn.

Glunder G

Klinik fur Geflugel, Tierarztlichen Hochschule Hannover.

Zentralblatt für Veterinarmedizin. Reihe B. Journal of veterinary medicine. Series B (GERMANY) Apr 1995, 42 (2) p89-99, ISSN 0514-7166 Journal Code: 0331325

Document type: Journal Article ; English Abstract

Languages: GERMAN

Main Citation Owner: NLM Record type: Completed Subfile: INDEX MEDICUS

Development of humoral precipitating antibodies against Campylobacter chickens . The development of precipitating antibodies in was examined by two-dimensional immunodiffusion test after chickens immunization with a formol inactivated vaccine and after subcutaneous and application of different live campylobacter serovars. The supernatant of bacterial cells after sonication and centrifugation was used as an antigen in the agar-gel precipitin test. Antisera against different serovars showed a high percentage of cross-reactions. In chickens immunized with an inactivated vaccine at an age of 1, 2, 3, 4 and 7 weeks, precipitating antibodies could be demonstrated for the first time at 7 days p.i. Except for 1-week-old birds, sera from the other groups reacted positively at 14 days p.o. After subcutaneous duplication of live organisms to 4-week-old chickens , antibodies could already be demonstrated at 4 days p.i. later in part of the experimental groups. No interrelation could be detected between antibody titers, measured by enzyme-linked immunoabsorbent assay (ELISA), from precipitating sera, as well as from those from non-precipitating sera. Precipitating antibodies and antibody titers in the ELISA were examined in sera from groups of birds infected at an age of 1, 2, 3, 4 and 7 weeks. During the Campylobacter excretion period, a distinct peak of antibody titers occurred in 1- and 7-week old birds, whereas other groups showed a steady increase in titers. Precipitating antibodies were only found in 1- and 2-week-old chickens.

Tags: Animal

Descriptors: Antibodies, Bacterial--biosynthesis--BI; *Bacterial Vaccines --immunology--IM; * Campylobacter --immunology--IM; * Campylobacter Infections--veterinary--VE; *Chickens; *Poultry Diseases--immunology--IM; Campylobacter Infections--immunology--IM; Immunodiffusion--veterinary--VE ; Vaccines, Inactivated -- immunology -- IM

CAS Registry No.: 0 (Antibodies, Bacterial); 0 (Bacterial Vaccines);

(Vaccines, Inactivated)

Record Date Created: 19951127 Record Date Completed: 19951127

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10200549
            96001607
                      PMID: 8526019
  Biological properties of yolk immunoglobulins.
  Janson A K; Smith C I; Hammarstrom L
  Dept of Clinical Immunology, Karolinska Institute, Huddinge Hospital,
Sweden.
  Advances in experimental medicine and biology (UNITED STATES)
                                                                          1995,
371A p685-90, ISSN 0065-2598 Journal Code: 0121103
  Document type: Journal Article
  Languages: ENGLISH
  Main Citation Owner: NLM
  Record type: Completed
  Subfile:
              INDEX MEDICUS
  Tags: Animal; Comparative Study; Female; Human
  Descriptors: Antibodies, Bacterial--immunology--IM; * Campylobacter
jejuni--immunology--IM; *Egg Proteins--immunology--IM; *Immunoglobulins
--immunology--IM; *Shigella flexneri--immunology--IM; Antibodies, Bacterial
--isolation and purification--IP; Bacterial Vaccines--immunology--IM;
Chickens; Egg Proteins--isolation and purification--IP; Enzyme-Linked
Immunosorbent
                  Assay;
                               Granulocytes--immunology--IM;
                                                                    Granulocytes
--microbiology--MI; Hela Cells; Hemagglutination Tests; Immunization,
Secondary; Immunoglobulins--isolation and purification--IP; Phagocytosis;
Shigella flexneri--physiology--PH; Vaccination
 CAS Registry No.: 0
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                                                           (Bacterial Vaccines);
    (Egg Proteins); 0
                         (IgY); 0
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 Record Date Created: 19960124
 Record Date Completed: 19960124
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06814168 91054093 PMID: 2241686

Influence of antibody treatment of Campylobacter jejuni on the dose required to colonize chicks.

Stern N J; Meinersmann R J; Dickerson H W

Poultry Microbiological Safety Research Unit, Richard B. Russell Agricultural Research Center, USDA-Agricultural Research Service, Athens, Georgia 30613.

Avian diseases (UNITED STATES) Jul-Sep 1990, 34 (3) p595-601, ISSN 0005-2086 Journal Code: 0370617

Document type: Journal Article

Languages: ENGLISH

Main Citation Owner: NLM Record type: Completed Subfile: INDEX MEDICUS

study was designed to clarify the role of controlling chicken colonization by Campylobacter jejuni. Cecal colonization by C. jejuni was compared after the organism was exposed either to phosphate-buffered saline, normal rabbit serum, rabbit hyperimmune anti-C. jejuni serum, or anti-C. jejuni antibodies extracted antibodies bile. Antibodies from chicken bile were extracted by from chicken affinity absorption against outer-membrane proteins from the challenge organism. Sera were heated 1 hour at 56 C to destroy complement activity. Bacterial inoculum levels were enumerated after 1 hour exposure at 4 C to the various treatments. The heated sera and the bile antibodies were not bactericidal, and bacterial agglutination was not evident. Serial dilutions of the antibody-treated C. jejuni were given by gavage into 1-day-old chicks. Six days later, the ceca were removed from the chicks, and samples were cultured Campylobacter -charcoal differential agar. The on colonization dose-50% was increased by twofold to 160-fold when the organism was preincubated with hyperimmune antiserum or the bile antibodies as compared with preincubation with phosphate-buffered saline. We conclude that antibodies inhibit chicken cecal colonization by C. jejuni.

Tags: Animal; Support, Non-U.S. Gov't; Support, U.S. Gov't, Non-P.H.S. Descriptors: Campylobacter Infections--veterinary--VE; * Campylobacter jejuni--immunology--IM; *Chickens; *Immunization, Passive; *Poultry Diseases--immunology--IM; Bile--immunology--IM; Campylobacter Infections--immunology--IM; Campylobacter jejuni--growth and development--GD; Carrier State--immunology--IM; Carrier State--veterinary--VE; Cecum --microbiology--MI; Colony Count, Microbial; Dose-Response Relationship, Immunologic; Enzyme-Linked Immunosorbent Assay; Immune Sera--immunology--IM; Immunoglobulin A, Secretory--immunology--IM; Immunoglobulin A, Secretory--isolation and purification--IP

CAS Registry No.: 0 (Immune Sera); 0 (Immunoglobulin A, Secretory) Record Date Created: 19901207

Record Date Created: 19901207 Record Date Completed: 19901207 06814167 91054092 PMID: 2241685

Influence of Campylobacter jejuni cecal colonization on immunoglobulin response in chickens .

Myszewski M A; Stern N J

Department of Medical Microbiology, University of Georgia, Athens 30606. Avian diseases (UNITED STATES) Jul-Sep 1990, 34 (3) p588-94, ISSN 0005-2086 Journal Code: 0370617

Document type: Journal Article

Languages: ENGLISH
Main Citation Owner: NLM
Record type: Completed
Subfile: INDEX MEDICUS

immunoglobulin response of chickens to colonization by jejuni isolates B-540 and Clin-1 was monitored. Chicken Campylobacter humoral IgG and biliary secretory IgA (sIgA) responses were assessed by enzyme-linked immunosorbent assay (ELISA). Samples were taken from 128 C. jejuni-colonized chickens and 104 uncolonized chickens housed in a controlled environment. An indirect ELISA was performed using the homologous isolate of C. jejuni as the capture antigen and was developed with the specific goat anti-chicken IgG or IgA alkaline phosphatase conjugates. The ELISA absorbance values of the test samples at 405 nm (serum diluted 1:32 and bile diluted 1:10) were normalized in direct proportion to standard sera and bile sample values. In the colonized chickens , humoral IgG activities were highest at hatch, dropped to their lowest level after 2 weeks, and increased by 8 weeks to levels similar to those detected at hatch. The sIgA activity was lowest at hatch and increased by 4 weeks in colonized chickens while remaining lower in the control chickens. Chickens colonized with isolate B-540 showed a primary sIgA response during the first 4 weeks and reached a plateau over the final 4 weeks. In spite of these limited humoral and secretory immunoglobulin responses, once the chicken ceca was colonized by C. jejuni, the organism persisted throughout the 8-week experiment. Tags: Animal

Descriptors: Campylobacter jejuni--immunology--IM; *Cecum--microbiology
--MI; * Chickens --immunology--IM; * Immunoglobulin A, Secretory
--biosynthesis--BI; *Immunoglobulin G--biosynthesis--BI; Bile--immunology
--IM; Campylobacter Infections--immunology--IM; Campylobacter
Infections--veterinary--VE; Campylobacter jejuni--growth and development
--GD; Carrier State--immunology--IM; Carrier State--veterinary--VE;
Enzyme-Linked Immunosorbent Assay; Poultry Diseases--immunology--IM
CAS Registry No.: 0 (Immunoglobulin A, Secretory); 0 (Immunoglobulin G)

Record Date Created: 19901207 Record Date Completed: 19901207 08383145 95071121 PMID: 7526839

Isotype, specificity, and kinetics of systemic and mucosal antibodies to Campylobacter jejuni antigens, including flagellin, during experimental oral infections of chickens.

Cawthraw S; Ayling R; Nuijten P; Wassenaar T; Newell D G

Central Veterinary Laboratory (Weybridge), New Haw, Addlestone, Surrey, United Kingdom.

Avian diseases (UNITED STATES) Apr-Jun 1994, 38 (2) p341-9, ISSN 0005-2086 Journal Code: 0370617

Document type: Journal Article

Languages: ENGLISH

Main Citation Owner: NLM Record type: Completed Subfile: INDEX MEDICUS

The **immune** response of **chickens** to **Campylobacter** jejuni infection was studied as a step in the search for vaccine candidates. One-day-old chicks orally challenged with C. jejuni strain 81116 showed significant increases in specific IgG, IgA, and IgM circulating **antibodies**, as detected by enzyme-linked immunosorbent assay (ELISA). These levels peaked at 9, 5, and 7 weeks postinfection, respectively. Maternal IgG antibodies were also detected over the first 2 weeks. Specific mucosal IgG and IgA levels also increased significantly. All of the antibody demonstrated a major response to the 62- kDa flagellin protein by Western blotting techniques. The immunodominance of flagellin was confirmed by ELISA using an antigen preparation from an aflagellate mutant. When overlapping recombinant polypeptide fragments of flagellin were used, epitopes detected by chicken antibodies were observed in region IV, between residues 95-340 of the protein. Thus flagellin may be suitable candidate for a vaccine, although its role in protection must first be established.

Tags: Animal; Support, Non-U.S. Gov't

Descriptors: Antibodies, Bacterial--biosynthesis--BI; *Antigens, Bacterial--immunology--IM; * Campylobacter Infections--immunology--IM; * Campylobacter jejuni--immunology--IM; *Flagellin--immunology--IM; *Immunoglobulin G--biosynthesis--BI; *Immunoglobulin Isotypes--biosynthesis--BI; *Intestinal Mucosa--immunology--IM; Antibodies, Bacterial--blood--BL; Antibodies, Bacterial--classification--CL; Antibody Specificity; Blotting, Western; Chickens; Enzyme-Linked Immunosorbent Assay; Epitopes--analysis --AN; Immunoglobulin G--blood--BL; Immunoglobulin G--classification--CL; Immunoglobulin Isotypes--classification--CL

CAS Registry No.: 0 (Antibodies, Bacterial); 0 (Antigens, Bacterial); 0 (Epitopes); 0 (Immunoglobulin G); 0 (Immunoglobulin Isotypes); 12777-81-0 (Flagellin)

Record Date Created: 19941129
Record Date Completed: 19941129

11732157 99169031 PMID: 10069862

Apoptotic ffect of outer-membrane proteins from Campylobacter jejuni on chicken lymphocytes.

Zhu J; Meinersmann R J; Hiett K L; Evans D L

Poultry Microbiological Safety Research Unit, Russell Agricultural Research Center, U.S. Department of Agriculture, Agricultural Research Service, Athens, GA 30604.

Current microbiology (UNITED STATES) Apr 1999, 38 (4) p244-9, ISSN 0343-8651 Journal Code: 7808448

Document type: Journal Article

Languages: ENGLISH

Main Citation Owner: NLM Record type: Completed

Subfile: INDEX MEDICUS; AIDS/HIV

jejuni is a significant cause of food-borne diseases in Campylobacter humans. The bacterium is considered a commensal organism in chickens , and it can heavily colonize chickens without causing inflammation. Poultry may be the major reservoir for the human infection in developed countries. Here we show that an outer-membrane protein extract prepared from the bacteria caused apoptosis of **chicken** lymphocytes detected in vitro with the terminal deoxynucleotidyl transferase-mediated dUTP nick end-labeling assay that preferentially labels individual apoptotic cells. Blood- and spleen-lymphocytes from different-aged chickens displayed a significantly greater percentage of apoptotic cells after culture with the outer-membrane proteins from C. jejuni than controls treated with phosphate-buffered saline, ovalbumin, or outer-membrane proteins prepared from E. chicken coli strain BL21. The C. jejuni extract also produced apoptosis of chicken lymphoblastoid tumor cell lines. Apoptosis was blocked by pretreating the extract with proteinase K or antiserum against outer-membrane proteins. The results suggest that C. jejuni may be capable of achieving immune avoidance in chickens by causing apoptosis of lymphocytes.

Tags: Animal; Support, U.S. Gov't, Non-P.H.S.

Descriptors: Apoptosis; *Bacterial Outer Membrane Proteins--pharmacology --PD; *CD4-Positive T-Lymphocytes--drug effects--DE; *CD8-Positive T-Lymphocytes--drug effects--DE; *Campylobacter jejuni--chemistry--CH; Age Factors; Bacterial Outer Membrane Proteins--drug effects--DE; *Campylobacter jejuni--immunology--IM; Chickens; Endopeptidase K--pharmacology--PD; Escherichia coli--chemistry--CH; Ovalbumin --pharmacology--PD; Specific Pathogen-Free Organisms; Tumor Cells, Cultured CAS Registry No.: 0 (Bacterial Outer Membrane Proteins); 9006-59-1 (Ovalbumin)

Enzyme No.: EC 3.4.21.64 (Endopeptidase K)

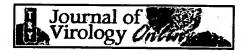
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           Biochemical Studies-Proteins, Peptides and Amino Acids
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           Biophysics-Molecular Properties and Macromolecules
   10508
           Biophysics-Membrane Phenomena
   12504
           Pathology, General and Miscellaneous-Diagnostic
           Blood, Blood-Forming Organs and Body Fluids-Blood and Lymph
   15002
              Studies
           Immunology and Immunochemistry-General; Methods
   34502
   36002
           Medical and Clinical Microbiology-Bacteriology
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           Veterinary Science-Microbiology
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           Biochemical Studies-Carbohydrates
           Medical and Clinical Microbiology-Serodiagnosis
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   85570 Trogoniformes
   85715 Bovidae
   85765 Canidae
   86215 Hominidae
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 09363587
           BIOSIS NO.: 199497371957
 Isotype, specificity, and kinetics of systemic and mucosal antibodies to
  Campylobacter jejuni antigens, including flagellin, during experimental
  oral infections of chickens.
AUTHOR: Cawthraw S; Ayling R; Nuijten P; Wassenaar T; Newell D G(a)
AUTHOR ADDRESS: (a) Central Vet. Lab., New Haw, Addlestone, Surrey KT15 3NB
JOURNAL: Avian Diseases 38 (2):p341-349 1994
ISSN: 0005-2086
DOCUMENT TYPE: Article
RECORD TYPE: Abstract
LANGUAGE: English
SUMMARY LANGUAGE: English; Spanish
ABSTRACT: The immune response of chickens to Campylobacter jejuni
  infection was studied as a step in the search for vaccine candidates.
  One-day-old chicks orally challenged with C. jejuni strain 81116 showed
  significant increases in specific IgG, IgA, and IgM circulating
  antibodies , as detected by enzyme-linked immunosorbent assay
  (ELISA). These levels peaked at 9, 5, and 7 weeks postinfection,
  respectively. Maternal IgG antibodies were also detected over the first
  2 weeks. Specific mucosal IgG and IgA antibody levels also increased
  significantly. All of the birds demonstrated a major response to the
  62- kDa flagellin protein by Western blotting techniques. The
  immunodominance of flagellin was confirmed by ELISA using an antigen
  preparation from an aflagellate mutant. When overlapping recombinant
  polypeptide fragments of flagellin were used, epitopes detected by
  chicken antibodies were observed in region IV, between residues 95-340
  of the protein. Thus flagellin may be a suitable candidate for a vaccine,
  although its role in protection must first be established.
DESCRIPTORS:
 MAJOR CONCEPTS: Dental and Oral System (Ingestion and Assimilation);
    Immune System (Chemical Coordination and Homeostasis); Infection;
   Pathology; Veterinary Medicine (Medical Sciences)
 BIOSYSTEMATIC NAMES: Aerobic Helical or Vibrioid Gram-Negatives--
   Eubacteria, Bacteria; Galliformes--Aves, Vertebrata, Chordata, Animalia
 ORGANISMS: aerobic helical or vibrioid gram-negative bacteria (Aerobic
   Helical or Vibrioid Gram-Negatives); Campylobacter jejuni (Aerobic
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Helical or Vibrioid Gram-Negatives); Galliformes (Galliformes BIOSYSTEMATIC CLASSIFICATION (SUPER TAXA): animals; bacteria; birds;

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chordates; eubacteria; microorganisms; nonhuman vertebrates;
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  MISCELLANEOUS TERMS:
                        PATHOGENICITY; VACCINATION
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          Dental and Oral Biology-Pathology
  19006
          Immunology and Immunochemistry-Bacterial, Viral and Fungal
  34504
  36002
          Medical and Clinical Microbiology-Bacteriology
          Veterinary Science-Pathology
          Veterinary Science-Microbiology
  38006
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         Biochemical Studies-Carbohydrates
BIOSYSTEMATIC CODES:
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          Galliformes
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   Isotype, specificity, and kinetics of systemic and mucosal antibodies to
 Campylobacter
                  jejuni antigens, including flagellin, during experimental
oral infections of chickens
  Cawthraw, S. Ayling, R.; Nuijten, P.; Wassenaar, T.; Newell, D.G.
  Kennett Square, Pa. : American Association of Avian Pathologists Inc.
  Avian diseases. Apr/June 1994. v. 38 (2) p. 341-349.
  ISSN: 0005-2086 CODEN: AVDIAI
  DNAL CALL NO: 41.8 Av5
  Language: English
                      Summary Language: Spanish
  Includes references
  Place of Publication: Pennsylvania
  Subfile: IND; OTHER US (NOT EXP STN, EXT, USDA; SINCE 12/76);
  Document Type: Article
        immune response of chickens to Campylobacter jejuni infection
  The
was studied as a step in the search for vaccine candidates. One-day-old
chicks orally challenged with C. jejuni strain 81116 showed significant
increases in specific IgG, IgA, and IgM circulating antibodies, as
detected by enzyme-linked immunosorbent assay (ELISA). These levels peaked
at 9, 5, and 7 weeks postinfection, respectively. Maternal IgG antibodies
 were also detected over the first 2 weeks. Specific mucosal IgG and IgA
 antibody
                    also increased significantly. All of the
            levels
 demonstrated a major response to the 62- kDa flagellin protein by Western
blotting techniques. The immunodominance of flagellin was confirmed by
ELISA using an antigen preparation from an aflagellate mutant. When
overlapping recombinant polypeptide fragments of flagellin were used,
epitopes detected by chicken antibodies were observed in region IV,
between residues 95-340 of the protein. Thus flagellin may be a suitable
candidate for a vaccine, although its role in protection must first be
established.
 DESCRIPTORS: chickens;
                           campylobacter jejuni; experimental infections;
 antibody formation;
                       iga; igg; igm; maternal antibodies; flagella;
bacterial antigens; antigenic determinants; isotypes;
  Section Headings: L832
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PHARMACOLOGY AND IMMUNE THERAPEUTIC AGENTS
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Institution: US PATENT & TRADEMARK OFFICE Sign In as Member

J. Bacteriol., Jun 1992, 3874-3883, Vol 174, No. 12 Copyright © 1992, American Society for Microbiology

Biochemical and antigenic properties of the Campylobacter flagellar hook protein

ME Power, RA Alm and TJ Trust

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The flagellar filament-hook complex was removed from Campylobacter cells by shearing and was purified by differential solubilization and ultracentrifugation at pH 11 followed by cesium chloride buoyant density ultracentrifugation. Flagellar filaments were then dissociated in 0.2 M glycine-HCl (pH 2.2), and purified hooks were collected by ultracentrifugation. The hooks (105 by 24 nm) each displayed a conical protrusion at the proximal end, a concave cavity at the distal end, and helically arranged subunits. The apparent subunit molecular weight of the hook protein of seven of the eight Campylobacter strains studied was 92,500, while that of the other was 94,000. N-terminal amino acid analysis of the hook protein of two strains of Campylobacter coli and one strain of Campylobacter jejuni demonstrated that the first 15 residues were identical. Amino acid composition analysis showed that the Campylobacter hook protein contained 35.7% hydrophobic and 9.5% basic residues. Isoelectric focusing determined that the hook protein was acidic, with a pI of 4.9. Comparisons with the Salmonella and Caulobacter hook protein compositions

Department of Biochemistry and Microbiology, University of Victoria, British Columbia, Canada.





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Institution: US PATENT & TRADEMARK OFFICE Sign In as Member

Infect. Immun., Oct 1994, 4256-4260, Vol 62, No. 10 Copyright © 1994, American Society for Microbiology

Heat shock- and alkaline pH-induced proteins of Campylobacter jejuni: characterization and immunological properties

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YL Wu, LH Lee, DM Rollins and WM Ching

Naval Medical Research Institute, Bethesda, Maryland 20889-5607.

The protein response to physiological stress was characterized in Campylobacter jejuni 81176 after exposure to heat and pH shock and following periods of recovery. Immunoreactivities of major stress- related proteins were determined with anti-Campylobacter immune rabbit serum and intestinal lavage fluid. Distinct proteins with molecular masses ranging from 10 to 120 kDa were induced and/or released by selective heat or pH treatments. The most notable responses were those of two proteins with apparent molecular masses of 45 and 64 kDa that were induced and two other proteins of 10 and 12 kDa that were released by selective heat shock, alkaline pH treatment, or both. On the basis of N-terminal sequence analysis and immunological cross-reactivity data, the 64- and 10-kDa proteins were the C. jejuni homologs of Escherichia coli GroEL and GroES proteins, respectively. Enhanced chemiluminescence Western blotting (immunoblotting) revealed that all four proteins were among the major protein antigens recognized by anti-

Entry name	Q9R4E6
Primary accession number	Q9R4E6
Secondary accession numbers	None
Entered in TrEMBL in	Release 13, May 2000
Sequence was last modified in	Release 13, May 2000
Annotations were last modified in	Release 25, September 2003

Name and origin of the protein	
Protein name	groEL-like stress protein 62 kDa subunit [Fragment]
Synonyms	None
Gene name	None
From	Campylobacter jejuni [TaxID: 197]
Taxonomy	Bacteria; Proteobacteria; Epsilonproteobacteria; Campylobacterales; Campylobacteraceae; Campylobacter.

References

[1] SEQUENCE.

MEDLINE=96123358; PubMed=8577276; [NCBI, ExPASy, EBI, Israel, Japan]

Takata T., Wai S.N., Takade A., Sawae Y., Ono J., Amako K.;

"The purification of a GroEL-like stress protein from aerobically adapted Campylobacter jejuni.";

Microbiol. Immunol. 39:639-645(1995).

Comments

None

Cross-references

0318986 96121222 PMID: 8576327

Identification and characterization of an immunogenic outer membrane protein of Campylobacter jejuni.

Burnens A; Stucki U; Nicolet J; Frey J

Institute for Veterinary Bacteriology, University of Berne, Switzerland. Journal of clinical microbiology (UNITED STATES) Nov 1995, 33 (11) p2826-32, ISSN 0095-1137 Journal Code: 7505564

Document type: Journal Article

Languages: ENGLISH Main Citation Owner: NLM Record type: Completed INDEX MEDICUS

We cloned and expressed in Escherichia coli a gene encoding an 18- kDa outer membrane protein (Omp18) from Campylobacter jejuni ATCC 29428. The nucleotide sequence of the gene encoding Omp18 was determined, and an open reading frame of 165 amino acids was revealed. The amino acid sequence had the typical features of a leader sequence and a signal peptidase II cleavage site at the N-terminal part of Omp18. Moreover, the sequence had a high degree of similarity to the peptidoglycan-associated outer membrane lipoprotein P6 of Haemophilus influenzae and the peptidoglycan-associated lipoprotein PAL of E. coli. Southern blot analysis in which the cloned gene was used as a probe revealed genes similar to that encoding Omp18 in all the thermophilic group of campylobacters as well as Campylobacter sputorum. All campylobacters tested expressed a protein with a molecular mass identical to that of Omp18. The protein reacted immunologically with **polyclonal antibodies** directed against Omp18 from C. jejuni. PCR amplification of the gene encoding Omp18 with specific primers and subsequent restriction enzyme analysis of the amplified DNA fragments showed that the gene for Omp18 is highly conserved in C. jejuni strains isolated from humans, dogs, cats, calves, and chickens but is different in other Campylobacter species. In order to obtain pure recombinant Omp18 protein for serological assays, the cloned gene for Omp18 was genetically modified by replacing the signal sequence with a DNA segment encoding six adjacent histidine residues. Expression of this in E. coli allowed purification of the modified protein (Omp18-6xHis) by metal chelation chromatography. Sera from patients with past C. jejuni infection reacted positively with Omp18-6xHis, while sera from healthy blood donors showed no reaction with this antigen. Omp18, which is an outer membrane protein belonging to the family of PALs is well conserved in C. jejuni and is highly immunogenic. It is therefore a good candidate as an antigen for the serological diagnosis of past C. jejuni infections.

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Entry information

069239
Q9PN75
Release 39, May 2000
Release 40, October 2001
Release 41, February 2003
60 kDa chaperonin
Protein Cpn60
groEL protein
GROL or GROEL or MOPA or CJ1221
Campylobacter jejuni [TaxID: 197]
Bacteria; Proteobacteria;
Epsilonproteobacteria;
Campylobacterales; Campylobacteraceae;
<u>Campylobacter</u> .

[1]|SEQUENCE FROM NUCLEIC ACID.

MEDLINE=99140140; PubMed=10206714; [<u>NCBI, ExPASy, EBI, Israel,</u> <u>Japan]</u>

Thies F.L., Weishaupt A., Karch H., Hartung H.P., Giegerich G.;

"Cloning, sequencing and molecular analysis of the Campylobacter jejunigroESL bicistronic operon.";

Microbiology 145:89-98(1999).

[2]|SEQUENCE FROM NUCLEIC ACID.

STRAIN=ATCC 43429, ATCC 43432, ATCC 43438, and ATCC 43456;

Cunningham A., Taboada E., Nash J.H., Wakarchuk W.W., Gilbert M.;

"Sequencing of the cpn60 gene from various Campylobacter jejuni isolates.";

Submitted (DEC-2001) to the EMBL/GenBank/DDBJ databases.

[3] SEQUENCE FROM NUCLEIC ACID.

STRAIN=NCTC 11168:

MEDLINE=20150912; PubMed=10688204; [NCBI, ExPASy, EBI, Israel, Japan]

Parkhill J., Wren B.W., Mungall K., Ketley J.M., Churcher C., Basham D.,

Chillingworth T., Davies R.M., Feltwell T., Holroyd S., Jagels K., Karlyshev

A.V., Moule S., Pallen M.J., Penn C.W., Quail M.A., Rajandream M.A.,

Rutherford K.M., van Vliet A.H.M., Whitehead S., Barrell B.G.;

"The genome sequence of the food-borne pathogen Campylobacter jejuni reveals hypervariable sequences.";

Nature 403:665-668(2000).

Comments

FUNCTION: Prevents misfolding and promotes the refolding and proper assembly of unfolded polypeptides generated under stress conditions (By similarity).

SUBUNIT: Oligomer of 14 subunits composed of two stacked rings of 7 subunits (By similarity).

SUBCELLULAR LOCATION: Cytoplasmic (By similarity).

SIMILARITY: Belongs to the chaperonin (HSP60) family.

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	4544044	[CoDingSequence]		
	AF461064;	[EMBL / GenBank / DDBJ]		
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	AAL67844.1;	[CoDingSequence]		
	AL139077;	[EMBL / GenBank / DDBJ]		
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PIR	G81328; G81328.			
HSSP	PO6139; 1GRL. [HSSP ENTRY / SWISS-3DIMAGE / PDB]			
CMR	069289; CJ1221.			
HAMAP	MF_00600; -; 1.			
7 17 17 17 11	PBIL [Family / Alignment / Tree]			
	IPRO01844; Chaprnin_Cpn60.			
InterPro	IPRO02423; Cpn60/TCP-1.			
	Graphical view of domain structure.			
Pfam	PF00118; cpn60_TCP1; 1.			
PRINTS	PRO0298; CHAPERONIN60.			
	PROO304; TCOMPLEXTCP1.			
PROSITE	PS00296; CHAPERONINS_CPN60; 1.			
ProDom	[Domain structure / List of seq. sharing at least 1 domain]			
HOBACGEN	[Family / Alignment / Tree]			
BLOCKS	069289.			
ProtoNet	<u>069289</u> .			
ProtoMap	<u>069289</u> .	·		

PRESAGE	<u>069289</u> .	
DIP	<u>069289</u> .	
ModBase	<u>069289</u> .	
SWISS-2DP	AGE Get region on 2D PAGE.	

Keywords

Chaperone; ATP-binding; Complete proteome.

Features



Feature table viewer

Key From To Length Description

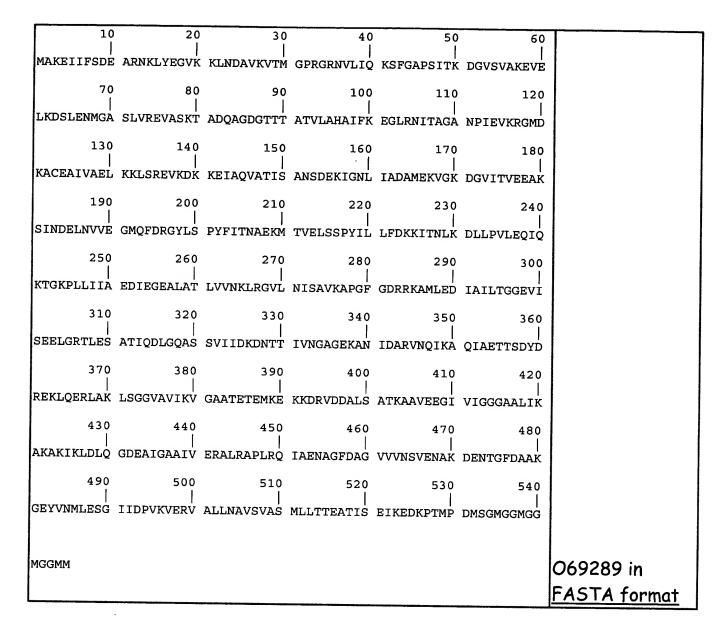
CONFLICT $\underline{179}$ $\underline{179}$ A -> P (IN REF. $\underline{1}$).

CONFLICT 383 383 A -> T (IN REF. $\underline{1}$).

Sequence information

Length: 545 Molecular weight: CRC64: 4DA80BC64330C41E [This is a

AA 57970 Da checksum on the sequence]



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BLAST submission on ExPASy/SIB or at NCBI (USA)



Sequence analysis tools: <u>ProtParam,</u>
<u>ProtScale, Compute pI/Mw,</u>
<u>PeptideMass, PeptideCutter,</u>
<u>Dotlet</u> (Java)



<u>ScanProsite,</u> <u>MotifScan</u>



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The Korean ExPASy site, kr.expasy.org, is temporarily not available.